Université de Liège Faculté des Sciences Département de Géologie Laboratoire de Minéralogie



# Pegmatite phosphates: from the field to the lab.

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- 1. Introduction
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Intro.

Fillowite + alluaudite, Kabira pegmatite, Uganda



Johnsomervilleite, Loch Quoich, Scotland



### Occurrence

-Granitic pegmatites -Metamorphic rocks -Meteorites

#### Chladniite, GRA 95209 meteorite



The Varuträsk pegmatite

Petro.



Varuträsk Skellefteå 5 km Paleoproterozoic (ca. 1.87 - 1.66 Ga) Paleoproterozoic (ca. 1.96 - 1.86 Ga) Revsund suite Acid to intermediate intrusive rocks Varuträsk formation Skellefte group Felsic to intermediate metavolcar Metabasic volcanic rocks (amphibolite, metabasalt) (granite, granodiorite) rocks (metarhyolite, metadacite, Bothnian / Vargfors group Metasedimentary rocks metaandesite) Paleoproterozoic (ca. 1.87 - 1.75 Ga) Metasedimentary carbonate rock Skellefte suite (calcitic to dolomitic marbles (metagreywackes, schists) Acid to intermediate intrusive rocks -Acid to intermediate intrusive rocks (granite, granodiorite) Deformation zone (granite, granodiorite) < Sunform - Anti

Field

Intro.



Stability



Crystallo.

Percy Quensel (1881-1966)

#### Brian Mason (1917-2009)











The alluaudite group

Petro.



Field

Intro.

#### Varulite, Na<sub>2</sub>Mn<sub>2</sub>Fe<sup>3+</sup>(PO<sub>4</sub>)<sub>3</sub> Varuträsk, Sweden

François II Alluaud (1778-1866) Mayor of Limoges and mineralogist Chanteloube pegmatite Alluaudite, NaMnFe<sup>3+</sup><sub>2</sub>(PO<sub>4</sub>)<sub>3</sub>



Augustin-Alexis Damour (1808-1902)





Crystallo. Sta

Stability

Crystallo. Field Petro.

Stability



## Genesis of alluaudites

Intro.



### **Oxidation mechanism**

 $Na_2MnFe^{2+}Fe^{3+}(PO_4)_3 \rightarrow []NaMnFe^{3+}_2(PO_4)_3$  $Na^+ + Fe^{2+} \implies [] + Fe^{3+}$ 

### Secondary origin Primary origin



Alluaudite, Kibingo pegmatite, Rwanda

Intro.

Petro.

Crystallo.

Stability

## Let's go to the field!









### Argentina



Crystallo.

Stability

### Pegmatite zoning

Field





#### MINERALOGY AND GEOCHEMISTRY OF PHOSPHATES AND SILICATES IN THE SAPUCAIA PEGMATITE, MINAS GERAIS, BRAZIL: GENETIC IMPLICATIONS

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<u>Fe-Mn phosphates in</u> <u>pegmatites</u>

Petro.

Field

Intro.

Crystallo.

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Palermo #1 pegmatite, NH



Buranga pegmatite, Rwanda



Intro.

Field

Petro.

Crystallo. Stability

### Back to the lab...



### Fe-Mn phosphates







### Petrography



Al phosphates

Thin sections

Intro.

Petro.

Crystallo. Stability

### <u>The triphylite + sarcopside</u> assemblage



#### Intercroissances et inclusions dans les associations graftonite-sarcopside-triphylite

par ANDRÉ-MATHIEU FRANSOLET, Institut de Minéralogie, Université de Liège (<sup>1</sup>).







Sarcopside (Fe,Mn)<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub>



<u>The alluaudite + fillowite</u> assemblage

Petro.

Field

Intro.



Crystallo.

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#### Alluaudite + fillowite, Kabira, Uganda



Stability

### The triphylite + alluaudite assemblage



#### PETROGRAPHIC EVIDENCE FOR PRIMARY HAGENDORFITE IN AN UNUSUAL ASSEMBLAGE OF PHOSPHATE MINERALS, **KIBINGO GRANITIC PEGMATITE, RWANDA**

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Hagendorfite, alluaudite, and heterosite, Kibingo pegmatite, Rwanda

### Complex assemblages from Sapucaia

Petro.

Intro.

Field





Crystallo.

Stability

Single-crystal X-ray diffraction

Petro.

Crystallo.

Stability





#### **4-circle diffractometer**

Field

Intro.



#### **Diffraction spots**



**Structure determination** 





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### NaFe<sup>2+</sup>PO<sub>4</sub>

#### a = 4.882(1), b = 10.387(2), c = 6.091(1) Å Pbnm



Karen Louise Webber



Malpensata pegmatite, Italy

Field

Stability

## <u>Zavalíaite, a new mineral...</u>



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#### a = 6.088(1) Å b = 4.814(1) Å c = 10.484(2) Å $\beta$ = 89.42(3)° S.G. P2<sub>1</sub>/c

### $Mn_3(PO_4)_2$



Florencia Márquez Zavalía





The alluaudite structure

Petro.

Field

Intro.

A(2)': gable disphenoidA(1): distorted cubeM(1): very distorted octahedronM(2): distorted octahedron



Crystallo.

Stability

 $[A(2)A(2)'][A(1)A(1)'A(1)''2]M(1)M(2)_2(PO_4)_3$ 



Crystal chemistry of natural alluaudites Moore & Ito (1979)

 $\begin{array}{l} A(2)' \Rightarrow \operatorname{Na^{+}}, \, {}^{\bullet} \operatorname{K^{+}} \\ A(1) \Rightarrow \operatorname{Na^{+}}, \, \operatorname{Mn^{2+}}, \, \operatorname{Ca^{2+}}, \, {}^{\bullet} \\ M(1) \Rightarrow \operatorname{Mn^{2+}}, \, \operatorname{Fe^{2+}}, \, \operatorname{Ca^{2+}}, \, \operatorname{Mg^{2+}} \\ M(2) \Rightarrow \operatorname{Fe^{3+}}, \, \operatorname{Fe^{2+}}, \, \operatorname{Mn^{2+}}, \, \operatorname{Mg^{2+}}, \, \operatorname{Li^{+}} \end{array}$ 

Fransolet et al. (1985, 1986, 2004)

**Oxidation mechanism:** 

Na<sup>+</sup> + Fe<sup>2+</sup>  $\Rightarrow$  • + Fe<sup>3+</sup>

 $Na_{2}MnFe^{2+}Fe^{3+}(PO_{4})_{3} \Rightarrow \bullet NaMnFe^{3+}_{2}(PO_{4})_{3}$  $Na_{2}Fe^{2+}_{2}Fe^{3+}(PO_{4})_{3} \Rightarrow \bullet NaFe^{2+}Fe^{3+}_{2}(PO_{4})_{3}$ 

Stability

### LIÈGE université

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To Christian Chopin, for 30 years of dedicated service to EJM

#### A new nomenclature scheme for the alluaudite supergroup

New nomenclature for alluaudites

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 $\frac{\text{Type 1: }^{M(2)}M^{2+} < 0.5}{\text{Na}M^{2+}\text{Fe}^{3+}_{2}(\text{PO}_{4})_{3}\text{: ALLUAUDITES}}$   $= \text{Na}M^{2+}\text{Mn}^{3+}_{2}(\text{PO}_{4})_{3}\text{: ROOT1}$   $\frac{\text{Type 2: } 0.5 < ^{M(2)}M^{2+} < 1.5}{\text{Na}_{2}M^{2+}\text{Fe}^{2+}\text{Fe}^{3+}(\text{PO}_{4})_{3}\text{: HAGENDORFITES}}$   $\text{Na}_{2}M^{2+}\text{Mn}^{2+}\text{Fe}^{3+}(\text{PO}_{4})_{3}\text{: VARULITES}}$   $\text{Na}_{2}M^{2+}\text{MgFe}^{3+}(\text{PO}_{4})_{3}\text{: ROOT2}$ 

<u>Hydrothermal experiments</u>

Petro.





**Opened gold capsules** 

Field

#### Hydrothermal lab

Intro.

#### **Gold tubes**

Stability

Crystallo.



#### Hydrothermal bomb

P = 1 kbar T = 400-800°C



<u>Stability of the triphylite +</u> sarcopside assemblage

Petro.

Field

Intro.





Crystallo.

Stability

 $LiFe^{2+}_{2.5}(PO_4)_2$ 

Decrease of the Li-content of triphylite, from 0.72 *a.p.f.u.* at 400°C, to 0.48 *a.p.f.u.* at 600°C
Increase of the Li-content of sarcopside, from 0.01 *a.p.f.u.* at 400°C, to 0.05 *a.p.f.u.* at 600°C
1-phase domain above 700°C Calculation of crystallisation temperatures for natural assemblages

800 700 - × Triphylite 600 Sacopside 500 -T (°C) 400 300 Triphylite + sarcopside 200 100 0 0.40 0.00 0.20 0.60 0.80 1.00 Li p.f.u.

Fe/(Fe+Mn) ratio of natural triphylites and sarcopsides close to 0.800

Phase diagram for the LiMn<sub>0.5</sub>Fe<sup>2+</sup><sub>2</sub>(PO<sub>4</sub>)<sub>3</sub> starting composition

Cañada 35 % sarcopside and 65 % triphylite T ~ 500°C

Tsoabismund 15 % sarcopside and 85 % triphylite T ~ 350°C



Intro.

Field





•  $\underline{Mn} \Rightarrow \text{fillowite } [NaMn_4(PO_4)_3]$ No maricite [NaFePO<sub>4</sub>] in pegmatites Varulite  $Na_2Mn_2Fe^{3+}(PO_4)_3$ 350-400°C

> **Hagendorfite**  $Na_2MnFe^{2+}Fe^{3+}(PO_4)_3$ 450-500°C

**Ferrohagendorfite**  $Na_2Fe^{2+}_2Fe^{3+}(PO_4)_3$ 550-600°C

Frédéric Hatert · André-Mathieu Fransolet · Walter V. Maresch

system

an experimental investigation of the Na2(Mn2-2xFe1+2x)(PO4)3



Ki = Kibingo, Rwanda

<u>The Na-in-triphylite</u> geothermometer

Field

Intro.

850 **Triphylite** Maricite 800 750 700 T (°C) 650 600 550 500 0.2 0.4 0.6 0.8 0

Petro.

Experimental investigation of the alluaudite + triphylite assemblage, and development of the Na-in-triphylite geothermometer: applications to natural pegmatite phosphates

Frederic Hatert · Luisa Ottolini · Peter Schmid-Beurmann



•In triphylite, Na can reach 0.08 *a.p.u.f.* at 800°C

•In maricite, Li can reach 0.10 *a.p.u.f.* at 700°C

•No partitioning below ca. 550°C

Geothermometer!



Stability





- Phosphates are « exotic » minerals, forming large masses in the most evolved parts of granitic pegmatites
- They are of great interest for pegmatologists, to:
- Understand pegmatite evolution during the post-magmatic stages (HT and LT hydrothermal, meteoric)
- ✓ Definine the T and oxygen fugacity conditions of pegmatites
- For mineralogists and solid-state scientists:
- They provide an infinite source of new mineral species
   Their exciting crystal structures are an inspiration for the development of new materials (alluaudites and triphylites in Li-ion batteries)